AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Original) A method of treating obesity in a mammal comprising administering to a mammal in need of such treatment at least one weight-loss promoting anticonvulsant and at least one compound that enhances the activity of norepinephrine and/or dopamine in amounts such that said treatment is effected.
- 2. (Original) The method according to claim 1 wherein said anticonvulsant is of the formula (I):

$$R_3$$
 R_4
 R_2
 R_3
 R_3
 R_3

wherein X is CH₂ or oxygen,

R₁ is hydrogen or alkyl,

 R_2 , R_3 , R_4 and R_5 are independently hydrogen or lower alkyl, and when X is CH_2 , R_4 and R_5 can be alkene groups joined to form a benzene ring and when X is oxygen, R_2 and R_3 and/or R_4 and R_5 together can be a methylenedioxy group of the following formula (II):

$$R_{1}$$

wherein

R₆ and R₇ are the same or different and are hydrogen, lower alkyl or are alkyl and are joined to form a cyclopentyl or cyclohexyl ring.

- 3. (Original) The method according to claim 2 wherein R_1 is hydrogen or a C_1 - C_4 alkyl, straight and branched chain, and R_2 , R_3 , R_4 , R_5 , R_6 and R_7 are a C_1 - C_3 alkyl, straight or branched chain.
- 4. (Original) The method according to claim 1 wherein said anticonvulsant is of the formula (III):

$$R_1$$
 \xrightarrow{Y} CH_2SO_2N $\xrightarrow{R_2}$

wherein R_1 is hydrogen or a halogen atom, R_2 and R_3 are the same or different and are each hydrogen or an alkyl having 1 to 3 carbon atoms, and one of X and Y is a carbon

atom and another is a nitrogen atom, provided that the group -CH₂SO₂NR₂R₃ is bonded to the carbon atom of either of X and Y, or an alkali metal salt thereof.

- 5. (Original) The method according to claim 1 wherein said anticonvulsant is zonisamide or topiramate.
- 6. (Original) The method according to claim 1 wherein said compound that enhances the activity of norepinephrine and/or dopamine effects said enhancement via uptake inhibition.
- 7. (Original) The method according to claim 1 wherein said compound that enhances the activity of norepinephrine and/or dopamine is bupropion, Atomoxetine or Reboxetine.
- 8. (Original) The method according to claim 1 wherein said anticonvulsant and said compound that enhances the activity of norepinephrine and/or dopamine are administered separately.
- 9. (Original) The method according to claim 1 wherein said anticonvulsant and said compound that enhances the activity of norepinephrine and/or dopamine are administered concurrently.

- 10. (Original) A method of reducing the risk of hypertension, diabetes or dyslipidaemia in a mammal comprising administering to a mammal in need of such reduction at least one weight-loss promoting anticonvulsant and at least one compound that enhances the activity of norepinephrine and/or dopamine in amounts such that said reduction is effected.
- 11. (Original) A method of treating obesity in mammal comprising administering to a mammal in need of such treatment a compound of formula (III):

$$R_1$$
 \xrightarrow{Y} CH_2SO_2N $\xrightarrow{R_2}$

wherein R₁ is hydrogen or a halogen atom, R₂ and R₃ are the same or different and are each hydrogen or an alkyl having 1 to 3 carbon atoms, and one of X and Y is a carbon atom and another is a nitrogen atom, provided that the group -CH₂SO₂NR₂R₃ is bonded to the carbon atom of either of X and Y, or an alkali metal salt thereof, in an amount sufficient to effect said treatment.

- 12. (Original) The method according to claim 11 wherein said compound is zonisamide.
- 13. (Original) A composition comprising at least one weight loss-promoting anticonvulsant and at least one compound that enhances the activity of norepinephrine and/or dopamine.
- 14. (Original) The composition according to claim 13 wherein said compound is in dosage unit form.
- 15. (Original) The composition according to claim 14 wherein said composition is in the form of a tablet or capsule.
- 16. (Original) The composition according to claim 13 wherein said anticonvulsant is zonisamide or topiramate.
- 17. (Original) The composition according to claim 13 wherein said compound that enhances the activity of norepinephrine and/or dopamine is bupropion.
- 18. (New) A method of reducing weight in an overweight subject, said method comprising: administering to an overweight subject a pharmaceutical composition

comprising zonisamide, in an amount effective to reduce weight in said subject, wherein said weight loss is significant and sustained.

- 19. (New) The method according to claim 18, wherein said effective amount of zonisamide is in the range of about 50 to about 1000 mg/day.
- 20. (New) The method according to claim 19, wherein said effective amount of zonisamide is in the range of about 100 to about 600 mg/day.
- 21. (New) The method according to claim 18, wherein said overweight subject is an obese subject.
- 22. (New) The method according to claim 18, wherein said pharmaceutical composition is administered by a route selected from the group consisting of oral, parenteral, topical, injection and rectal administration.
- 23. (New) The method according to claim 22, wherein said pharmaceutical composition is administered orally to said subject.

- 24. (New) The method according to claim 18, wherein said pharmaceutical composition is administered in combination with another therapeutic method commonly used to treat overweight.
- 25. (New) The method according to claim 24, wherein said pharmaceutical composition is administered in combination with a hypocaloric diet or exercise.
- 26. (New) The method according to claim 24, wherein said pharmaceutical composition is administered in combination with orlistat, phentermine, sibutramine, topiramate, or sibutramine hydrochloride.
- 27. (New) A method of treating eating disorders in a subject in need of such treatment, said method comprising: administering to a subject a pharmaceutical composition comprising zonisamide, in an amount effective to treat eating disorders.
- 28. (New) The method according to claim 27, wherein said eating disorders are binge-eating disorder, bulimia nervosa, or anorexia nervosa.
- 29. (New) The method according to claim 27, wherein said effective amount of zonisamide is in the range of about 50 to about 1000 mg/day.

- 30. (New) The method according to claim 29, wherein said effective amount of zonisamide is in the range of about 100 to about 600 mg/day.
- 31. (New) The method according to claim 27, wherein said pharmaceutical composition is administered by a route selected from the group consisting of oral, parenteral, topical, injection and rectal administration.
- 32. (New) The method according to claim 31, wherein said pharmaceutical composition is administered orally to said subject.
- 33. (New) The method according to claim 27, wherein said pharmaceutical composition is administered in combination with another therapeutic agent commonly used to treat eating disorders.
- 34. (New) The method according to claim 33, wherein said therapeutic agent is fluoxetine, topiramate, or orlistat.
- 35. (New) A method of reducing weight in an overweight subject, said method comprising: administering to an overweight subject a pharmaceutical composition comprising zonisamide in an amount effective to induce weight loss in said subject, wherein the induction of weight loss is sustained during the dosing regimen.

- 36. (New) The method according to claim 35, wherein said effective amount of zonisamide is in the range of about 50 to about 1000 mg/day.
- 37. (New) The method according to claim 36, wherein said effective amount of zonisamide is in the range of about 100 to about 600 mg/day.
- 38. (New) The method according to claim 35, wherein said overweight subject is an obese subject.
- 39. (New) The method according to claim 35, wherein said pharmaceutical composition is administered by a route selected from the group consisting of oral, parenteral, topical, injection and rectal administration.
- 40. (New) The method according to claim 39, wherein said pharmaceutical composition is administered orally to said subject.
- 41. (New) The method according to claim 35, wherein said pharmaceutical composition is administered in combination with another therapeutic method commonly used to treat overweight.

- 42. (New) The method according to claim 41, wherein said pharmaceutical composition is administered in combination with a hypocaloric diet or exercise.
- 43. (New) The method according to claim 42, wherein said pharmaceutical composition is administered in combination with orlistat, phentermine, sibutramine, topiramate, or sibutramine hydrochloride.